

Trilotab 10 mg chewable tablets for dogs

Authorised

- Trilostane

Product identification

Medicine name:

Trilotab 10 mg chewable tablets for dogs

Active substance:

Trilostane

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Trilostane

10.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

Package description:

Aluminium-PVC/Aluminium/oPA blisters, containing 10 tablets. Cardboard box of 30 tablets

Aluminium-PVC/Aluminium/oPA blisters, containing 10 tablets. Cardboard box 100 tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:

CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:

25/08/2023

Manufacturing sites for batch release:

CP-Pharma Handelsgesellschaft mbH

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 129610

Date of authorisation status change:

25/08/2023

Reference member state:

Netherlands

Procedure number:

NL/V/0373/001

Concerned member states:

Austria Belgium Czechia Denmark Estonia Finland France Germany
Hungary Ireland Italy Latvia Lithuania Poland Portugal Slovakia Spain
Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

Published on: 22/08/2023

Updated on: 23/08/2023

[Download](#)

PuAR Trilotab NL_V_0373_001-005_DC 2023-08.pdf